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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
10/651,136	08/28/2003	Sandor Sipka	22740-2 8175 EXAMINER	
24256 7	590 04/24/2006			
DINSMORE & SHOHL, LLP 1900 CHEMED CENTER		NOLAN, PATRICK J		
255 EAST FIF			ART UNIT	PAPER NUMBER
CINCINNATI, OH 45202		1644		

DATE MAILED: 04/24/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

					
		Applicat	ion No.	Applicant(s)	
Office Action Summary		10/651,1	136	SIPKA ET AL.	
		Examine	or	Art Unit	
	·	Patrick J	. Nolan	1644	
Period fo	The MAILING DATE of this communicate Reply	ation appears on th	ne cover sheet with the c	orrespondence address	
A SH WHIC - Exter after - If NC - Failu Any (ORTENED STATUTORY PERIOD FOR CHEVER IS LONGER, FROM THE MAI notions of time may be available under the provisions of SIX (6) MONTHS from the mailing date of this commun of period for reply is specified above, the maximum statute to reply within the set or extended period for reply will reply received by the Office later than three months after the patent term adjustment. See 37 CFR 1.704(b).	LING DATE OF T 37 CFR 1.136(a). In no e ication. ory period will apply and v I, by statute, cause the ap	HIS COMMUNICATION vent, however, may a reply be tinwill expire SIX (6) MONTHS from plication to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).	
Status			•		
· ·	Responsive to communication(s) filed This action is FINAL . 2b Since this application is in condition for closed in accordance with the practice)⊠ This action is r allowance excep	non-final. t for formal matters, pro		
Dispositi	on of Claims				
5)□ 6)⊠ 7)□ 8)□ Applicati	Claim(s) 1-21 is/are pending in the app 4a) Of the above claim(s) 6-9,11,12,14 Claim(s) is/are allowed. Claim(s) 1-5,10,13 and 17-19 is/are re Claim(s) is/are objected to. Claim(s) are subject to restriction on Papers The specification is objected to by the second or	is/a jected. on and/or election Examiner. i) accepted or bon to the drawing(s)	requirement.	Examiner. e 37 CFR 1.85(a).	
11)	The oath or declaration is objected to b	y the Examiner. N	lote the attached Office	Action or form PTO-152.	
Priority ι	ınder 35 U.S.C. § 119				
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
2) 🔲 Notic 3) 🔯 Inforr	t(s) e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTC mation Disclosure Statement(s) (PTO-1449 or PT r No(s)/Mail Date <u>10-04-04</u> .		4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:		

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1. Claims 1-21 are pending.

2. Applicant's election of species, human and route of administration, aerosol in the reply

filed on 2-6-06 is acknowledged. Because applicant did not distinctly and specifically point out

the supposed errors in the restriction requirement, the election has been treated as an election

without traverse (MPEP § 818.03(a)).

3. Upon further consideration by the Examiner, claims 6 and 16, reading upon topical

administration does not read upon the elected invention, aerosol.

4. Claims 6-9, 11-12, 14-16, 20 and 21 are withdrawn from further consideration pursuant

to 37 CFR 1.142(b) as being drawn to a nonelected species, there being no allowable generic or

linking claim. Election was made without traverse in the reply filed on 2-06-06.

5. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making

and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode

contemplated by the inventor of carrying out his invention.

6. Claims 1-5, 10, 13 and 17-19 are rejected under 35 U.S.C. 112, first paragraph, as failing

to comply with the written description requirement. The claim(s) contains subject matter which

was not described in the specification in such a way as to reasonably convey to one skilled in the

relevant art that the inventor(s), at the time the application was filed, had possession of the

claimed invention.

Applicant has no written support in the specification as filed for the genus term LPS

derived from microbial, protozoan and/or fungal endotoxin.

The scope of the term reads upon thousands if not millions of LPS molecules isolated

from organism not yet even isolated. In a review of the specification, Applcant has written

support for LPS derived from E-coli bacteria. There is no disclosure of the additional LPS

structures from the varied genus Applicant is claiming.

Applicant is invited to review MPEP 2163

For each claim drawn to a genus:

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The written description requirement for a claimed genus may be satisfied through sufficient description of a representative number of species by actual reduction to practice(see i)(A), above), reduction to drawings (see i)(B), above), or by disclosure of relevant, identifying characteristics, i.e., structure or other physical and/or chemical properties, by functional characteristics coupled with a known or disclosed correlation between function and structure, or by a combination of such identifying characteristics, sufficient to show the applicant was in possession of the claimed genus (see i)(C), above). See Eli Lilly,119 F.3d at 1568, 43 USPQ2d at 1406.

A "representative number of species" means that the species which are adequately described are representative of the entire genus. Thus, when there is substantial variation within the genus, one must describe a sufficient variety of species to reflect the variation within the genus. >The disclosure of only one species encompassed within a genus adequately describes a claim directed to that genus only if the disclosure "indicates that the patentee has invented species sufficient to constitute the gen[us]." See Enzo Biochem, 323 F.3d at 966, 63 USPQ2d at 1615; Noelle v. Lederman, 355 F.3d 1343, 1350, 69 USPQ2d 1508, 1514 (Fed. Cir. 2004) (Fed. Cir. 2004)("[A] patentee of a biotechnological invention cannot necessarily claim a genus after only describing a limited number of species because there may be unpredictability in the results obtained from species other than those specifically enumerated."). "A patentee will not be deemed to have invented species sufficient to constitute the genus by virtue of having disclosed a single species when ... the evidence indicates ordinary artisans could not predict the operability in the invention of any species other than the one disclosed." In re Curtis, 354 F.3d 1347, 1358, 69 USPQ2d 1274, 1282 (Fed. Cir. 2004)(Claims directed to PTFE dental floss with a frictionenhancing coating were not supported by a disclosure of a microcrystalline wax coating where there was no evidence in the disclosure or anywhere else in the record showing applicant conveyed that any other coating was suitable for a PTFE dental floss.)< On the other hand, there may be situations where one species adequately supports a genus. See, e.g., Rasmussen, 650 F.2d at 1214, 211 USPQ at 326-27 (disclosure of a single method of adheringly applying one layer to another was sufficient to support a generic claim to "adheringly applying" because one skilled in Application/Control Number: 10/651,136

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the art reading the specification would understand that it is unimportant how the layers are adhered, so long as they are adhered); In re Herschler, 591 F.2d 693, 697, 200 USPQ 711, 714 (CCPA 1979) (disclosure of corticosteroid in DMSO sufficient to support claims drawn to a method of using a mixture of a "physiologically active steroid" and DMSO because "use of known chemical compounds in a manner auxiliary to the invention must have a corresponding written description only so specific as to lead one having ordinary skill in the art to that class of compounds. Occasionally, a functional recitation of those known compounds in the specification may be sufficient as that description."); In re Smythe, 480 F.2d 1376, 1383, 178 USPQ 279, 285 (CCPA 1973) (the phrase "air or other gas which is inert to the liquid" was sufficient to support a claim to "inert fluid media" because the description of the properties and functions of the air or other gas segmentizing medium would suggest to a person skilled in the art that appellant's invention includes the use of "inert fluid" broadly.). However, in Tronzo v. Biomet, 156 F.3d at 1159, 47 USPQ2d at1833 (Fed. Cir. 1998), the disclosure of a species in the parent application did not suffice to provide written description support for the genus in the child application. What constitutes a "representative number" is an inverse function of the skill and knowledge in the art. Satisfactory disclosure of a "representative number" depends on whether one of skill in the art would recognize that the applicant was in possession of the necessary common attributes or features of the elements possessed by the members of the genus in view of the species disclosed. For inventions in an unpredictable art, adequate written description of a genus which embraces widely variant species cannot be achieved by disclosing only one species within the genus. See, e.g., Eli Lilly. Description of a representative number of species does not require the description to be of such specificity that it would provide individual support for each species that the genus embraces. For example, in the molecular biology arts, if an applicant disclosed an amino acid sequence, it would be unnecessary to provide an explicit disclosure of nucleic acid sequences that encoded the amino acid sequence. Since the genetic code is widely known, a disclosure of an amino acid sequence would provide sufficient information such that one would accept that an applicant was in possession of the full genus of nucleic acids encoding a given amino acid sequence, but not necessarily any particular species. Cf. In re Bell, 991 F.2d 781, 785, 26 USPQ2d 1529, 1532 (Fed. Cir. 1993) and In re Baird, 16 F.3d 380, 382, 29 USPQ2d 1550, 1552 (Fed. Cir. 1994). If a representative number of adequately described species are not disclosed for

a genus, the claim to that genus must be rejected as lacking adequate written description under 35 U.S.C. 112, 1st.

7. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1-5, 10, 17-19 rejected under 35 U.S.C. 103(a) as being unpatentable over Tulic 8. et al.(reference U on the PTO-892), in view Matricardi et al., (Reference AO on the IDS submitted 10-4-04) and Bertok et al (Reference AN on the IDS submitted 10-4-04).

Tulic et al., teaches the prevention of allergy in 10 day old, immature mice administered LPS in aerosol, where said administration occurs weekly and daily and shortly after birth.

The claimed invention differs from the prior art teachings by the administration irradiated LPS to said mammals.

However, Matricardi et al., specifically teaches the administration of LPS has been shown to be beneficial to treat allergy, however a less toxic derivative of LPS would be preferred for treatment purposes. Bertok et al., teaches making irradiated LPS according to claim 2, and that it is a less toxic form of LPS that still stimulates an immune response. While the reference is Application/Control Number: 10/651,136

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silent to the type of immune response, based upon its cytokine profile, it would now be considered a Th1 immune resoponse.

Therefore, one of ordinary skill in the art at the time the invention was made would have been motivated to treat the mammals of Tulic et al, with an irradiated LPS molecules taught by Bertok et al., because said irradiated LPS molecule is loess toxic but maintains its immunostimulatory properties which is preferred as taught by Matricardi et al.

9. Claims 1 and 13 rejected under 35 U.S.C. 103(a) as being unpatentable over Tulic et al., in view of Matricardi et al., and Bertok et al., as applied to claims 1-5, 10, 17-19 above, and further in view of Liu et al.

Tulic et al., Matricardi et al., and Bertok et al., have been discussed supra.

The claimed invention differs from the prior art teachings by the recitation of treating a human of 1 month to 2 years of age.

However, Liu et al., specifically teaches that there ample data demonstrating that early life, less than two years, exposure to endotoxins, such as LPS, in humans has been demonstrated to decrease allergic sensitization and that frequent benign exposures to endotoxin early in life should be expected to influence immune development to prevent atopy, allergic disease and asthma.

Therefore one of ordinary skill in the art at the time the invention was made would have been motivated to use the treatment method taught by the combination of Tulic et al., in view of Matricardi et al., and Bertok et al. and treat young humans since Liu et al., recognizes early exposure to LPS in a benign way should be effective in preventing allergy.

The fax number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR

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system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Patrick Nolan whose telephone number is 571-272-0847.

If attempts to reach the examiner are unsuccessful, the examiner's supervisor, Christina Chan, can be reached at 571-272-0841.

Patrick J. Nolan, Ph.D.

Primary Examiner, Group 1640

April 14, 2006